## UK Patent Application (19) GB (11) 2 111 560 A

- (21) Application No 8229523
- (22) Date of filing 15 Oct 1982
- (30) Priority data
- (31) 330242
- (32) 14 Dec 1981
- (33) United States of America
  (US)
- (43) Application published 6 Jul 1983
- (51) INT CL<sup>3</sup> CO9K 7/O2
- (52) Domestic classification E1F GP
- (56) Documents cited
- (58) Field of search E1F
- (71) Applicant
  NL Industries Inc.,
  (USA—New Jersey),
  1230 Avenue of the
  Americas,
  New York,
  New York 10020,
  United States of America
- (72) Inventors

  Roy Francis House,

  Lonnie Daniel Hoover
- (74) Agent and/or address for service
  Michael Burnside and Partners,
  2 Serjeants Inn,
  Fleet Street,
  London,
  EC4Y 1HL

## (54) Viscous heavy brines

(57) Hydroxyethyl ceilulose and a sequestrant are added to a heavy brine containing one or more salts selected from calcium chloride, calcium bromide, and zinc bromide to increase the viscosity of the brine. Preferably the brine contains zinc bromide, has a density in the range from 1.70 to 2.30 g/cm³, and the sequestrant is a polyphosphonic acid, e.g. nitrilotri (methylenephosphonic acid) or water soluble salt thereof.

## SPECIFICATION Viscous heavy brines

The present invention relates to viscous brine solutions and, more particularly, to so called heavy brines having a density greater than 1.62 g/cm<sup>3</sup>. In recent years, the practical operating range of clear brines for use in the oil and gas industry has been significantly extended by utilizing soluble zinc salts, particularly zinc bromide, so that the advantages of clear brines can now be obtained with fluids having densities as high as 2.30 g/cm3 at ambient temperatures and pressures. The high density clear brines are used extensively: as completion fluids, to minimize plugging of 10 perforation tunnels, to protect formation permeability, and to minimize mechanical problems; as 10 workover fluids, for the same reasons; as packer fluids, to allow easy movement and retrieval of the packer; or under-reaming, gravel-pack and sand consolidation applications; as kill fluid on ballast fluid; for wire-line work; and as drilling fluids. Clear brines having a density of up to 1.70 g/cm3 are generally formulated to contain sodium 15 chloride, sodium bromide, potassium chloride, calcium chloride, calcium bromide or mixtures of these 15 salts. Clear brines having a density up to 1.81 g/cm3 can be formulated with calcium chloride and calcium bromide. If the brine must have a low crystallization temperature, however, clear brines in this density range are generally formulated to contain a soluble zinc salt. Zinc bromide is preferred because brines containing it are less corrosive than brines containing zinc chloride. Clear brines having a density 20 greater than 1.81 g/cm³ are formulated to contain zinc bromide. Viscous clear fluids are sometimes desired. Generally, hydroxyethyl cellulose (HEC) and xanthan gum polymers are compatible with the fluids which do not contain zinc salts. At higher densities, however, the hydration of the viscosifiers is significantly slower. HEC is generally considered as unsatisfactory for use in fluids containing zinc salts. It is an object of the present invention to provide viscous heavy brine solutions, particularly such 25 25 solutions containing zinc bromide and HEC. Another object of this invention is to provide a method of increasing the rate of hydration of HEC in heavy brines, particularly those brines having a density greater than 1.62 g/cm<sup>3</sup>. Still another object of this invention is to provide a method of increasing the viscosity of a heavy 30 brine by adding thereto an "activated" HEC, as disclosed more fully hereinafter, and a sequestrant. 30 The above and other objects of the present invention will become more apparent from the description given herein and the appended claims. This invention provides a method of increasing the viscosity of a heavy brine solution containing one or more soluble salts selected from calcium chloride, calcium bromide, zinc bromide, and mixtures 35 thereof, which comprises adding to said solution hydroxyethyl cellulose and from 0.25 to 20 g/l of 35 sequestrant. This invention also provides a method of increasing the rate of hydration of hydroxyethyl cellulose in a heavy brine solution containing one or more soluble salts selected from calcium chloride, calcium bromide, zinc bromide and mixtures thereof, which comprises adding to said brine from 0.25 to 20 g/l 40 40 of a sequestrant. This invention further provides a viscous heavy brine solution containing hydroxyethyl cellulose, one or more soluble salts selected from calcium chloride, calcium bromide, zinc bromide, and mixtures thereof, and from 0.25 to 20 g/l of a sequestrant. The present invention is based on the discovery that the addition of sequestrants to heavy brine 45 solutions increases the rate of hydration of HEC in the heavy brine and increases the viscosity of the 45 heavy brine containing HEC. The heavy brines for use in the present invention contain two or more soluble salts selected from calcium chloride, calcium bromide, zinc bromide, and mixtures thereof. Brines containing only calcium chloride can be formulated having a density from 1.02 to 1.40 g/cm3. Brines containing only calcium 50 bromide can be formulated having a density from 1.02 to 1.70 g/cm3. However, because calcium 50 chloride is much less expensive than calcium bromide, brines in the density range from 1.35 to 1.81 g/cm³ are generally formulated to contain both calcium chloride and calcium bromide, depending on the lowest temperature at which the brine will be used. Brines heavier than 1.80 g/cm<sup>3</sup> are generally formulated to contain calcium chloride, calcium bromide, and zinc bromide, or only calcium bromide 55 and zinc bromide, depending on the lowest temperature at which the brines will be used. Brines in the 55 density range from 1.70 to 1.80 g/cm3 may be formulated to contain calcium chloride, calcium bromide, and zinc bromide, if brines having a lower crystallization point are desired.

Generally, brines of any density within the range disclosed are prepared by mixing together various standard, commercially available brines, as follows: calcium chloride brines having a density from 1.32 to 1.39 g/cm³; calcium bromide brine having a density of 1.70 g/cm³; and a calcium bromide/zinc bromide brine having a density of 2.30 g/cm³ containing about 20% by weight of calcium

in the present invention. As indicated, such brines will generally have a density from 1.70 to 2.30

The most difficult brines to viscosify contain zinc bromide, and these brines are preferred for use

10

15

20

25

30

35

55

60

55

bromide and about 57% by weight of zinc bromide. Solid anhydrous calcium chloride and solid calcium bromide are also used in conjunction with these brines to prepare the heavy brines for use in this invention. Standard brine mixing tables are available from the various manufacturers of these brines.

The HEC polymers which are useful as viscosifiers in the present invention are solid, particulate materials which are water soluble or water dispersible and which upon solution or dispersion in an aqueous medium increase the viscosity of the system. HEC polymers are generally high yield, water soluble, non-ionic materials produced by treating cellulose with sodium hydroxide followed by reaction with ethylene oxide. Each anhydroglucose unit in the cellulose molecule has three reactive hydroxy groups. The average number of moles of the ethylene oxide that becomes attached to each anhydroglucose unit in cellulose is called moles of substituent combined. In general, the greater the degree of substitution, the greater the water solubility. In general, it is preferable to use HEC polymers having as high a mole substitution level as possible.

Usually, upon the addition of dry, powdered hydrophilic materials, such as HEC, to water, the polymer particles undergo hydration preventing the interior of the particle from readily hydrating, solvating or otherwise dispersing in the aqueous medium. Accordingly, high shear long mixing times and/or elevated temperatures must be applied in order to obtain a homogeneous system.

We have found that HEC and other hydrophilic polymers can be activated such that the polymers will viscosify heavy brines at ambient temperatures. Activated HEC compositions, and methods for activating HEC, are disclosed in our prior British Patent Applications No. 8101828 (Publication No. 2070611 A) and No. 8109880 (Publication No. 2075041 A). Methods of activating other hydrophilic polymers are disclosed in our prior British Patent Application No. 8125937 (Publication No. 2086923 A).

Activated HEC compositions comprise: (1) HEC, a solvating agent comprising a water miscible, polar organic liquid which when uniformly mixed with HEC in a weight ratio of HEC to solvating agent of 1:2 produces a mixture with substantially no free liquid solvating agent present after remaining quiescent for one week at ambient temperature in a sealed container, and a diluting agent comprising an organic liquid which is not a solvating agent; and (2) HEC, a water soluble organic liquid, and an aqueous liquid, the organic liquid when uniformly mixed with the HEC in a weight ratio of HEC to organic liquid of 1:2 produces a mixture with free liquid present after remaining quiescent for one week at ambient temperature in a sealed container, the aqueous liquid having a pH greater than 7.0.

Generally speaking, it has been found that virtually any organic compound which passes the solvation test described above, will function, to a usable degree, as a solvation agent. Non-limiting but preferred solvating agents include: aliphatic glycols containing from 2 to 5 carbon atoms, such as ethylene glycol, 1,2-propanediol, 1,4-butanediol, and 1,3-pentanediol; alkylene triols containing from 2 to 6 carbon atoms such as glycerol, 1,2,3-butanetriol and 1,2,3-pentanetriol; amides containing from 1 to 4 carbon atoms such as formamide, acetamide, and dimethyl formamide; and mixtures of the various above compounds.

The diluting agent, in general, will be any liquid organic compound or material which is not a solvating agent. In general, the diluting agents are liquids which do not appreciably swell the HEC 40 polymers, i.e. they do not produce semi-solid or viscous mixtures which have no free liquid present 40 after the one week solvation period described in the above test for determining solvating agents. Nonlimiting examples of diluting agents include liquid aliphatic and aromatic hydrocarbons containing from 5 to 10 carbon atoms, kerosene, diesel oil, isopropanol, alkylene glycol ethers, and vegetable oils. Particularly preferred are organic liquids which are water soluble or miscible, most preferably alkanols 45 having at least 3 carbon atoms, ethylene glycol monoalkyl ethers, and dialkylene glycol monoalkyl 45 ethers. The diluting agent will maintain the polymeric composition in a liquid, pourable state at a temperature of about 20°C. It will be understood, however, that lesser amounts of diluting agent can be used if desired, and that the ultimate amount of diluting agent employed will depend upon the type of shear which is available to disperse the thickener. In general it has been found that desirable thickeners, which are pourable liquids, can be produced from compositions containing from 10 to 25% 50 by weight of HEC polymer, from 2 to 70% by weight of diluting agent, and from 5 to 88% by weight of solvating agent.

The concentration of HEC need only be a viscosity increasing amount. Generally, the heavy brines of this invention will contain 0.25 to 15 g/l, preferably 0.7 to 8.6 g/l of HEC.

The sequestrants which are useful in the present invention include polybasic acids, e.g. polyphosphonic acids, aminopolycarboxylic acids, or polyhydroxy carboxylic acids, and their water soluble salts, particularly the alkali metal, ammonium, and alkanol ammonium salts thereof.

Representative, non-limiting, aminopolycarboxylic acids are compounds of the formula

## R<sub>1</sub>R<sub>2</sub>N[(CH<sub>2</sub>)<sub>8</sub>—NR<sub>3</sub>]<sub>6</sub>R<sub>4</sub>

60 where R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, and R<sub>4</sub> are independently H, —CH<sub>2</sub>COOH, or —(CH<sub>2</sub>CH<sub>2</sub>O)<sub>c</sub>H, c is 1 to 5, a is 2 to 6, and b is 0 to 6, provided that at least one-half of the radicals represented by R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, and R<sub>4</sub> are —CH<sub>2</sub>COOH.

Representative, non-limiting, polyhydroxycarboxylic acids are citric acid and gluconic acid.

Representative, non-limiting, polyphosphonic acids are set forth in U.S. Patent No. 3,971,734. Preferred polyphosphonic acids are free acids and water soluble salts of the formula

where each R is independently hydrogen, — $CH_2PO(OH)_2$ , or — $(C_nH_{2n}O)_cH$  wherein n is 2 to 4 and c is 5 1 to 3;  $R_1$  is

X | ---C---PO(OH)<sub>2</sub> | Y

wherein X and Y are independently hydrogen or  $C_1$ — $C_4$  alkyl, a is 2 to 6 and b is 0 to 6; provided that at least 1/2 of the radicals represented by R are — $CH_2PO(OH)_2$ ;

10 where R is —CH<sub>2</sub>PO(OH)<sub>2</sub>, R<sub>1</sub> is hydrogen, hydroxyl, or —(C<sub>n</sub>H<sub>2n</sub>O)<sub>c</sub>H wherein n is 2 to 4 and c is 1 to 3, 10 and where R<sub>2</sub> is  $\hat{C}_1$ — $\hat{C}_5$  alkyl or a group of the formula

wherein a is 0 to 8; or

where R is —CH<sub>2</sub>PO(OH)<sub>2</sub>, each R<sub>1</sub> is independently hydrogen, —CH<sub>2</sub>PO(OH)<sub>2</sub>, or (C<sub>n</sub>H<sub>2n</sub>O)<sub>c</sub>H, wherein 15 n is 2 to 4 and c is 1 to 3, and where R<sub>2</sub> is C<sub>1</sub>—C<sub>5</sub> alkyl, and

wherein a is 0 to 8.

Most particularly preferred stabilizing compounds are

$$\begin{array}{c|c}
R \\
\hline
R \\
1
\end{array}$$

$$\begin{array}{c|c}
CH_2 \\
a \\
R
\end{array}$$

$$\begin{array}{c|c}
D \\
R
\end{array}$$

$$\begin{array}{c|c}
D \\
D \\
D
\end{array}$$

20

where R is  $-CH_2PO(OH)_2$ ,  $R_1$  is  $-CH_2PO(OH)_2$ , a is either 2 or 6 and b is 0 to 4, especially those compounds wherein b is 0 or wherein a is 6 and b is 1 to 4.

Other water soluble polyphosphonates that contain at least two phosphonic acid groups per molecule which may be used in the practice of this invention are disclosed in U.S. Patents No. 25 3,733,270; 3,674,804; 3,576,783; 3,556,762; 3,551,339; 3,549,728; 3,497,313; 3,440,148; and 3,400,176.

The concentration of sequestrant useful in the practice of this invention will be an amount

sufficient to increase the rate of hydration of HEC in the heavy brine which is desired to be viscosified. Generally, concentrations of sequestrant from 0.25 to 20 g/l are sufficient.

To Illustrate the present invention more fully, the following non-limiting Example is presented. All physical property measurements were made in accordance with testing procedures set forth in Standard Procedure for Testing Drilling Fluid, API RP 13B, Seventh Edition, April, 1978. The physical property parameters referred to in the Example are in units expressed as follows:

Apparent viscosity=centipoise.

Plastic viscosity=centipoise.

Yield Point=Kg/m².

10 10-Sec. Gel Strength=Kg/m<sup>2</sup>.

10

15

20

25

55

Example of the invention

An activated HEC gellant was prepared by mixing together 124.5 parts by weight of isopropyl alcohol, 0.5 parts by weight of CAB-O-SIL M5 fumed silica, 50.0 parts by weight of NATROSOL 250 HHR hydroxyethyl cellulose, and 75.0 parts by weight of ethylene glycol. A 1.86 g/cm³ CaBr₂/ZnBr₂ solution was prepared by mixing a 1.70 g/cm³ CaBr₂ solution with a 2.30 g/cm³ CaBr₂/ZnBr₂ solution in a 0.74/0.26 volume ratio. Other 1.86 g/cm³ CaBr₂/ZnBr₂ solutions were prepared similarly, except that a 50% by weight aqueous solution of nitrilotri (methylenephosphonic acid) was added in varying amounts to the 2.30 g/cm³ solution before its dilution with the 1.70 g/cm³ solution. The concentration of nitrilotri (methylenephosphonic acid) in the 1.86 g/cm³ brines were 2.40, 5.89, and 20.74 g/l as indicated in Table 1. Thereafter, the brines were viscosified with 5.72 g/l of HEC (28.6 g/l of the activated gellant composition) by mixing with a Multimixer for 15 minutes. Fann viscosities were then obtained, after the solutions had hydrated for 90 hours at room temperature, and after the solutions were hot rolled for 16 hours at 65.5°C. The data obtained are given in Table 1.

The data indicate that the sequestrant increased the rate of hydration of the HEC and significantly increased the viscosity of the viscosified brine. However, the brine containing 20.74 g/l of this sequestrant had poor thermal stability as evidenced by the decrease in viscosity on rolling at 65.5°C.

Table I 5.72 g/I HEC g/I Nitrilotri (methylenephosphonic acid)

30 Fann V-G Rheology	3,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				
	0	2.40	5.89	20.74	30
After 15 Minutes Mixing				400	
600 rpm					
300 rpm					
Apparent Viscosity (cp)					35
Plastic Viscosity (cp)					33
Yield Point (Kg/m²)					
10-Sec. Gel Strength (Kg/m²)	0.05	0.15	0.20	0.24	
After 90 Hours Hydration					
600 rpm					4.0
300 rpm					40
Apparent Viscosity (cp)					
Plastic Viscosity (cp)					
Yield Point (Kg/m²)					
10-Sec. Gel Strength (Kg/m²)	0.20	0.54	1.51	2.05	
After Rolling Overnight at 65.5°C					45
	177	239			
•	124	164			
	88	119			
	53	75			
1 2	3.47	4.34			50
10-Sec. Gel Strength (Kg/m²)	0.68	1.17	1.71	0.07	
% Hydration in 15 Minutes*	31.5	40.9	46.7	115.9	
	After 15 Minutes Mixing 600 rpm 300 rpm Apparent Viscosity (cp) Plastic Viscosity (cp) Yield Point (Kg/m²) 10-Sec. Gel Strength (Kg/m²)  After 90 Hours Hydration 600 rpm 300 rpm Apparent Viscosity (cp) Plastic Viscosity (cp) Yield Point (Kg/m²) 10-Sec. Gel Strength (Kg/m²) 10-Sec. Gel Strength (Kg/m²) After Rolling Overnight at 65.5°C 600 rpm 300 rpm Apparent Viscosity (cp) Plastic Viscosity (cp) Plastic Viscosity (cp) Yield Point (Kg/m²)	### After 15 Minutes Mixing  600 rpm  39  Apparent Viscosity (cp)  Plastic Viscosity (cp)  Yield Point (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  Apparent Viscosity (cp)  After 90 Hours Hydration  600 rpm  300 rpm  48  Apparent Viscosity (cp)  Plastic Viscosity (cp)  Yield Point (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  After Rolling Overnight at 65.5°C  600 rpm  300 rpm  177  300 rpm  Apparent Viscosity (cp)  Plastic Viscosity (cp)  After Rolling Overnight at 65.5°C  6100 rpm  177  300 rpm  124  Apparent Viscosity (cp)  Plastic Viscosity (cp)  Plastic Viscosity (cp)  S3  Yield Point (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  3.47  10-Sec. Gel Strength (Kg/m²)  0.68	### After 15 Minutes Mixing  600 rpm  300 rpm  39  Apparent Viscosity (cp)  Plastic Viscosity (cp)  Yield Point (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  Apparent Viscosity (cp)  600 rpm  600 rpm  600 rpm  88  121  Apparent Viscosity (cp)  Plastic Viscosity (cp)  Plastic Viscosity (cp)  Yield Point (Kg/m²)  1.95  2.78  10-Sec. Gel Strength (Kg/m²)  After Rolling Overnight at 65.5°C  600 rpm  124  Apparent Viscosity (cp)  After Rolling Overnight at 65.5°C  600 rpm  124  Apparent Viscosity (cp)  Plastic Viscosity (cp)  88  119  Plastic Viscosity (cp)  78  300 rpm  124  Apparent Viscosity (cp)  53  75  Yield Point (Kg/m²)  3.47  4.34  10-Sec. Gel Strength (Kg/m²)  0.68  1.17	### After 15 Minutes Mixing  600 rpm  600 rpm  39 67 86  Apparent Viscosity (cp)  Plastic Viscosity (cp)  23 32 39  Yield Point (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  Plastic Viscosity (cp)  78 92 120  Plastic Viscosity (cp)  78 92 120  Plastic Viscosity (cp)  Plastic Viscosity (cp)  78 92 120  Plastic Viscosity (cp)  Plastic Viscosity (cp)  78 92 120  Plastic Viscosity (cp)  88 119 123  Plastic Viscosity (cp)  75 75 62  Yield Point (Kg/m²)  75 62  Yield Point (Kg/m²)  76 75 62  Yield Point (Kg/m²)  77 1.71	### After 15 Minutes Mixing  600 rpm  39 67 86 95  Apparent Viscosity (cp)  Plastic Viscosity (cp)  23 32 39 43  Yield Point (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  Plastic Viscosity (cp)  31 185 240 280  300 rpm  39 67 86 95  10-Sec. Gel Strength (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  10-Sec. Gel Strength (Kg/m²)  11-Sec. Gel Strength (Kg/m²)  12-Sec. Gel Strength (Kg/m²)  13-Sec. Gel Strength (Kg/m²)

300 rpm Reading After 15 Minutes×100

Claims

<sup>\*%</sup> Hydration in 15 Minutes=\_\_\_\_\_\_\_300 rpm Reading After Rolling @ 65.5°C

<sup>1.</sup> A method of increasing the viscosity of a heavy brine solution containing one or more soluble

	salts selected from calcium chloride, calcium bromide, zinc bromide, and mixtures thereof, which comprises adding to said solution hydroxyethyl cellulose and from 0.25 to 20 g/l of sequestrant.  2. A method as claimed in Claim 1, wherein the hydroxyethyl cellulose is added to the brine as an	
5	activated gellant composition in which the hydroxyethyl cellulose has been activated to hydrate in heavy brines at ambient temperatures.	5
	3. A method as claimed in Claim 1 or 2, wherein the concentration of hydroxyethyl cellulose is from 0.25 to 15 g/l.	
	4. A method as claimed in any of the preceding claims, wherein the sequestrant is a	
	polyphosphonic acid, aminopolycarboxylic acid, polyhydroxy carboxylic acid, or a water soluble salt	10
10	thereof.	10
	5. A method as claimed in Claim 4, wherein the sequestrant is nitrilotri (methylenephosphonic acid) or a water soluble salt thereof.	
	6. A method of increasing the rate of hydration of hydroxyethyl cellulose in a heavy brine solution	
	containing one or more soluble salts selected from calcium chloride, calcium bromide, zinc bromide	
15	and mixtures thereof, which comprises adding to said brine from 0.25 to 20 g/l of a sequestrant.	15
13	7. A method as claimed in Claim 6, wherein the brine contains from 0.25 to 15 g/l of	. •
	hydroxyethyl cellulose.	
	8. A method as claimed in Claim 6 or 7, wherein the sequestrant is a polyphosphonic acid,	
	aminopolycarboxylic acid, polyhydroxy carboxylic acid, or a water soluble salt thereof.	
20	9. A method as claimed in Claim 8, wherein the sequestrant is nitrilotri (methylenephosphonic	20
	acid) or a water soluble salt thereof.	
	10. A viscous heavy brine solution containing hydroxyethyl cellulose, one or more soluble salts	
	selected from calcium chloride, calcium bromide, zinc bromide, and mixtures thereof, and from 0.25 to	
	20 g/l of a sequestrant.	
25	11. A viscous heavy brine as claimed in Claim 10 which comprises from 0.25 to 15 g/l of	25
	hydroxyethyl cellulose.	
	12. A viscous heavy brine as claimed in Claim 10 or 11, wherein the sequestrant is a	
	polyphosphonic acid, aminopolycarboxylic acid, polyhydroxy carboxylic acid, or a water soluble salt	
	thereof.	20
30	13. A viscous heavy brine as claimed in Claim 12, wherein the sequestrant is nitrilotri	30
	(methylenephosphonic acid) or a water soluble salt thereof.	
	14. A method as claimed in Claim 1 or 6 and substantially as hereinbefore described with reference to the Example.	
	15. A viscous heavy brine as claimed in Claim 10 and substantially as hereinbefore described	
35	with reference to the Example.	35
55	With following to the Extension	-